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APPLICATION NO.	FILING DATE	FIRST NAMED INVENTOR	ATTORNEY DOCKET NO.	CONFIRMATION NO.
10/697,716	10/31/2003	H. William Bosch	029318-0977	8372
	7590 12/15/200 very, Inc. c/o Foley & I	EXAMINER		
3000 K Street, I		JEAN-LOUIS, SAMIRA JM		
Suite 500 Washington, DC 20007-5109			ART UNIT	PAPER NUMBER
<i>C</i> ,			1617	
			MAIL DATE	DELIVERY MODE
			12/15/2008	PAPER

Please find below and/or attached an Office communication concerning this application or proceeding.

The time period for reply, if any, is set in the attached communication.

Advisory Action Before the Filing of an Appeal Brief

Application No.	Applicant(s)	
10/697,716	BOSCH ET AL.	
Examiner	Art Unit	

	SAMIRA JEAN-LOUIS	1617	
The MAILING DATE of this communication appe	ars on the cover sheet with the c	correspondence add	ress
THE REPLY FILED 20 November 2008 FAILS TO PLACE THIS	S APPLICATION IN CONDITION F	OR ALLOWANCE.	
1. The reply was filed after a final rejection, but prior to or on application, applicant must timely file one of the following application in condition for allowance; (2) a Notice of Apperfor Continued Examination (RCE) in compliance with 37 Comperiods:	replies: (1) an amendment, affidavit eal (with appeal fee) in compliance	t, or other evidence, w with 37 CFR 41.31; or	hich places the (3) a Request
a) The period for reply expires 3 months from the mailing date b) The period for reply expires on: (1) the mailing date of this A no event, however, will the statutory period for reply expire to Examiner Note: If box 1 is checked, check either box (a) or (dvisory Action, or (2) the date set forth i ater than SIX MONTHS from the mailing	date of the final rejection	in.
MONTHS OF THE FINAL REJECTION, See MPEP 706.07(f).		
Extensions of time may be obtained under 37 CFR 1.136(a). The date whave been filed is the date for purposes of determining the period of extunder 37 CFR 1.17(a) is calculated from: (1) the expiration date of the set forth in (b) above, if checked. Any reply received by the Office later may reduce any earned patent term adjustment. See 37 CFR 1.704(b). NOTICE OF APPEAL	ension and the corresponding amount on hortened statutory period for reply origing than three months after the mailing date	of the fee. The appropria nally set in the final Offic	ate extension fee e action; or (2) as
 The Notice of Appeal was filed on A brief in comp filing the Notice of Appeal (37 CFR 41.37(a)), or any exter Notice of Appeal has been filed, any reply must be filed with AMENDMENTS 	nsion thereof (37 CFR 41.37(e)), to	avoid dismissal of the	
3. ☐ The proposed amendment(s) filed after a final rejection, b	out prior to the date of filing a brief,	will not be entered be	cause
(a) They raise new issues that would require further cor	nsideration and/or search (see NOT		
(c) ☐ They are not deemed to place the application in bet appeal; and/or	·	ducing or simplifying tl	ne issues for
(d) ☐ They present additional claims without canceling a c NOTE: (See 37 CFR 1.116 and 41.33(a)).	corresponding number of finally reje	ected claims.	
4. The amendments are not in compliance with 37 CFR 1.12			PTOL-324).
 Applicant's reply has overcome the following rejection(s): 			
 Newly proposed or amended claim(s) would be all non-allowable claim(s). 	owable if submitted in a separate, t	imely filed amendmer	nt canceling the
7. For purposes of appeal, the proposed amendment(s): a) [how the new or amended claims would be rejected is prov The status of the claim(s) is (or will be) as follows: Claim(s) allowed: Claim(s) objected to: Claim(s) rejected:		l be entered and an e:	xplanation of
Claim(s) withdrawn from consideration:			
AFFIDAVIT OR OTHER EVIDENCE			
 The affidavit or other evidence filed after a final action, but because applicant failed to provide a showing of good and was not earlier presented. See 37 CFR 1.116(e). 			
 The affidavit or other evidence filed after the date of filing entered because the affidavit or other evidence failed to o showing a good and sufficient reasons why it is necessary 	vercome <u>all</u> rejections under appea and was not earlier presented. Se	ll and/or appellant fail ee 37 CFR 41.33(d)(1	s to provide a).
10. ☐ The affidavit or other evidence is entered. An explanation REQUEST FOR RECONSIDERATION/OTHER	n of the status of the claims after er	ntry is below or attach	ed.
11. The request for reconsideration has been considered but See Continuation Sheet.	t does NOT place the application in	condition for allowan	ce because:
12. Note the attached Information <i>Disclosure Statement</i> (s). (13. Other:	PTO/SB/08) Paper No(s)		
/SREENI_PADMANABHAN/ Supervisory Patent Examiner, Art Unit 1617			

Continuation Sheet (PTO-303)

Application No.

The Examiner respectfully points out that in the response to Amendment in the Final rejection dated 09/16/08, the rejection under 35 U.S.C.102 (b) was withdrawn. Subsequently, a rejection of claims 1-5, 7, 9-14, 18-21, 28-41, and 43-47 was made under 35 U.S.C.103 (a). Furthermore, given that the co-pending application 10/683,154 has been abandoned, the ODP rejection in the Final Office Action dated 09/16/08 is hereby withdrawn.

Applicant's traversal of the 35 U.S.C.103 (a) rejection of claims 1-5, 7, 9-14, 18-21, 28-41, and 43-47 over Krause in view Radhakrishnan has been fully considered but is not found persuasive. Given that Krause teaches triamcinalone that are encapsulated within PLA particles and given that Krasue also teach PLA particle sizes of less than 1 micron, the Examiner contends that one of ordinary skill in the art would have concluded that the triamcinalone cannot be bigger than the capsule that contains it. As a result, one of ordinary skill in the art would have envisaged that the PLA encapsulated triamcinalone particles would necessarily have to be of a size less than 1 micron. As for applicant's arguments that Krause does not specify the effective triamcinalone particle size, such arguments are again not persuasive as Krause clearly teaches that a suspension of the nanoparticles (PLA encapsulated triamcinalone) was maintained for each distribution wherein the size of the PLA particles were from 500 nm, 476 nm and 710 nm; all of which are under 1 micron (see pg. 147, Size distribution and table 1). Thus, Krause does indeed render obvious applicant's invention.

Applicant's traversal of the PLA as a stabilizer or a capsule has been fully considered but is not found persuasive. The Examiner reiterates the fact that the drug is embedded in the center of the PLA sphere which necessarily mean that the PLA particle is adsorbed unto (i.e. on top of) the triamcinalone and this reads on applicant's limitation. Moreover, the Examiner respectfully points out that the PLA can also serve more than one purpose in a solution. In this instance, it can prevent agglomeration of particles due to encapsulation of the drug and second, as a stabilizer that allows the small triamcinalone particles to disperse effectively. Additionally, Krause teaches the addition of other stabilizers to his composition such as gelatin for emulsification purposes (see pg. 147, Preparation of PLA particles) thereby suggesting to one of ordinary skill in the art to add gelatin if an emulsified composition is desired. Thus, given that applicant did not define a stabilizer as solely a surfactant, the Examiner contends that anything that stabilizes the compound such as PLA can necessarily be viewed as a stabilizer. Thus, for the foregoing reasons, the Examiner contends that Krause does indeed render obvious applicant's invention.

Appplicant's traversal of Unger has again been fully considered but is not found persuasive. Unger was provided to demonstrate that encapsulated drugs including triamcinalone, aspirin, and emulsifiers can all be encapsulated and administered topically. Therefore, Unger does indeed render obvious applicant's claims 6, 17, and 22. For the foregoing reasons, the Examiner contends that the 103 (a) rejections over Krause in view of Radhakrishan and in further view of Unger were indeed proper and are MAINTAINED.